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NEWS 2	APR 04	STN AnaVist, Version 1, to be discontinued
NEWS 3	APR 15	WPIDS, WPINDEX, and WPIX enhanced with new predefined hit display formats
NEWS 4	APR 28	EMBASE Controlled Term thesaurus enhanced
NEWS 5	APR 28	IMSRESEARCH reloaded with enhancements
NEWS 6	MAY 30	INPAFAMDB now available on STN for patent family searching
NEWS 7	MAY 30	DGENE, PCTGEN, and USGENE enhanced with new homology sequence search option
NEWS 8	JUN 06	EPFULL enhanced with 260,000 English abstracts
NEWS 9	JUN 06	KOREAPAT updated with 41,000 documents
NEWS 10	JUN 13	USPATFULL and USPAT2 updated with 11-character patent numbers for U.S. applications
NEWS 11	JUN 19	CAS REGISTRY includes selected substances from web-based collections
NEWS 12	JUN 25	CA/CAplus and USPAT databases updated with IPC reclassification data
NEWS 13	JUN 30	AEROSPACE enhanced with more than 1 million U.S. patent records
NEWS 14	JUN 30	EMBASE, EMBAL, and LEMBASE updated with additional options to display authors and affiliated organizations
NEWS 15	JUN 30	STN on the Web enhanced with new STN AnaVist Assistant and BLAST plug-in
NEWS 16	JUN 30	STN AnaVist enhanced with database content from EPFULL
NEWS 17	JUL 28	CA/CAplus patent coverage enhanced
NEWS 18	JUL 28	EPFULL enhanced with additional legal status information from the epoline Register
NEWS 19	JUL 28	IFICDB, IFIPAT, and IFIUDB reloaded with enhancements
NEWS 20	JUL 28	STN Viewer performance improved
NEWS 21	AUG 01	INPADOCDB and INPAFAMDB coverage enhanced
NEWS 22	AUG 13	CA/CAplus enhanced with printed Chemical Abstracts page images from 1967-1998
NEWS 23	AUG 15	CAOLD to be discontinued on December 31, 2008
NEWS 24	AUG 15	CAplus currency for Korean patents enhanced
NEWS 25	AUG 25	CA/CAplus, CASREACT, and IFI and USPAT databases enhanced for more flexible patent number searching
NEWS 26	AUG 27	CAS definition of basic patents expanded to ensure comprehensive access to substance and sequence information
NEWS 27	SEP 18	Support for STN Express, Versions 6.01 and earlier, to be discontinued
NEWS 28	SEP 25	CA/CAplus current-awareness alert options enhanced to accommodate supplemental CAS indexing of exemplified prophetic substances

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 NEWS 29  SEP 26  WPIDS, WPINDEX, and WPIX coverage of Chinese and
                  and Korean patents enhanced
 NEWS 30  SEP 29  IFICLS enhanced with new super search field
 NEWS 31  SEP 29  EMBASE and EMBAL enhanced with new search and
                  display fields
 NEWS 32  SEP 30  CAS patent coverage enhanced to include exemplified
                  prophetic substances identified in new Japanese-
                  language patents

 NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,
              AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

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FILE 'HOME' ENTERED AT 14:47:41 ON 06 OCT 2008

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COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
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STRUCTURE FILE UPDATES:    5 OCT 2008  HIGHEST RN 1057399-47-9
DICTIONARY FILE UPDATES:   5 OCT 2008  HIGHEST RN 1057399-47-9

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=> s cloforabine
L1            0 CLOFORABINE
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=> s clofarabine
L2            2 CLOFARABINE

=> d l2 1-2

L2   ANSWER 1 OF 2  REGISTRY  COPYRIGHT 2008 ACS on STN
RN   134646-41-6  REGISTRY
ED   Entered STN:  05 Jul 1991
CN   9H-Purin-6-amine, 2-chloro-9-[2-deoxy-2-fluoro-5-O-
     [hydroxy[[hydroxy(phosphonooxy)phosphinyl]oxy]phosphinyl]-β-D-
     arabinofuranosyl]-  (CA INDEX NAME)
OTHER NAMES:
CN   Clofarabine triphosphate
FS   STEREOSEARCH
MF   C10 H14 Cl F N5 O12 P3
SR   CA
LC   STN Files:   CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.
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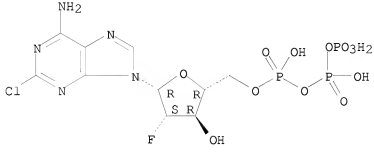


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**PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**

             17 REFERENCES IN FILE CA (1907 TO DATE)
             17 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L2   ANSWER 2 OF 2  REGISTRY  COPYRIGHT 2008 ACS on STN
RN   123318-82-1  REGISTRY
ED   Entered STN:  20 Oct 1989
CN   9H-Purin-6-amine, 2-chloro-9-(2-deoxy-2-fluoro-β-D-arabinofuranosyl)-
     (CA INDEX NAME)
OTHER NAMES:
CN   2-Chloro-9-(2-deoxy-2-fluoro-β-D-arabinofuranosyl)adenine
CN   Clofarabine
CN   Clofarex
CN   Clolar
FS   STEREOSEARCH
MF   C10 H11 Cl F N5 O3
CI   COM
SR   CA
LC   STN Files:   ADISINSIGHT, ANABSTR, BEILSTEIN*, BIOSIS, CA, CAPLUS,
       CASREACT, CBNB, CHEMCATS, CIN, CSCHEM, EMBASE, IMSDRUGNEWS, IMSPATENTS,
       IMSPRODUCT, IMSRESEARCH, IPA, MEDLINE, PATDPASPC, PHAR, PROMT, PROUSDDR,
       RTECS*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL
         (*File contains numerically searchable property data)

Absolute stereochemistry.
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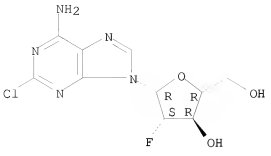

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

170 REFERENCES IN FILE CA (1907 TO DATE)
5 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
171 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> file caplus
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	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	15.22	15.43

FILE 'CAPLUS' ENTERED AT 14:49:11 ON 06 OCT 2008
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FILE COVERS 1907 - 6 Oct 2008 VOL 149 ISS 15
FILE LAST UPDATED: 5 Oct 2008 (20081005/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

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http://www.cas.org/legal/infopolicy.html

=> s 123318-82-1/rn
171 123318-82-1
5 123318-82-1D
L3 169 123318-82-1/RN
(123318-82-1 (NOTL) 123318-82-1D)

=> s l3 and lupus
21128 LUPUS
L4 3 L3 AND LUPUS

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=> d l4 1-3 ibib abs hitstr

L4   ANSWER 1 OF 3  CAPLUS  COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:         2007:1086774  CAPLUS
DOCUMENT NUMBER:          147:413153
TITLE:                    Indole derivatives as inhibitors of histone
                          deacetylase
INVENTOR(S):              Buggy, Joseph J.; Balasubramanian, Sriram; Verner,
                          Erick; Tai, Vincent W.-F.; Lee, Chang-Sun
PATENT ASSIGNEE(S):       Pharmacyclics, Inc., USA
SOURCE:                   PCT Int. Appl., 137pp.
                          CODEN: PIXXD2
DOCUMENT TYPE:            Patent
LANGUAGE:                 English
FAMILY ACC. NUM. COUNT:   1
PATENT INFORMATION:

     PATENT NO.          KIND   DATE        APPLICATION NO.        DATE
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     WO 2007109178        A2    20070927    WO 2007-US6714         20070316
     WO 2007109178        A3    20080731
         W:  AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
             GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN,
             KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, MG, MK, MN,
             MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS,
             RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ,
             UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
         RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
             IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW,
             GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
             BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA
     US 20070281934        A1    20071206    US 2007-687565         20070316
PRIORITY APPLN. INFO.:                     US 2006-783287P     P   20060316
OTHER SOURCE(S):          MARPAT 147:413153
AB   Described herein are compds. and pharmaceutical compns. containing such
     compds., which inhibit the activity of histone deacetylase 8 (HDAC8).
     Also described herein are methods of using such HDAC8 inhibitors, alone
     and in combination with other compds., for treating diseases or conditions
     that would benefit from inhibition of HDAC8 activity.  The synthesis of
     some of the active compds. is described.  Tablets contained a HDAC8
     inhibitor at 400 mg/tablet.
IT   123318-82-1, Clolar
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (indole derivs. as inhibitors of histone deacetylase)
RN   123318-82-1  CAPLUS
CN   9H-Purin-6-amine, 2-chloro-9-(2-deoxy-2-fluoro-β-D-arabinofuranosyl)-
       (CA INDEX NAME)

Absolute stereochemistry.
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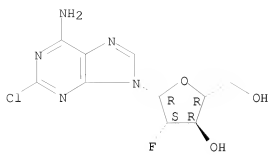


L4 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:619578 CAPLUS
DOCUMENT NUMBER: 147:46112
TITLE: Treatment of cancer and other diseases
INVENTOR(S): Habib, Nabil
PATENT ASSIGNEE(S): Nabil Habib Lab, Lebanon; Vianova Labs, Inc.
SOURCE: PCT Int. Appl., 86pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007064691	A1	20070607	WO 2006-US45665	20061130
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
CA 2632903	A1	20070607	CA 2006-2632903	20061130
EP 1968607	A1	20080917	EP 2006-844623	20061130
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				

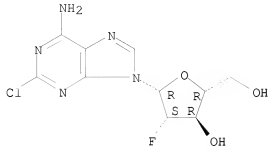
PRIORITY APPLN. INFO.: US 2005-741725P P 20051202
WO 2006-US45665 W 20061130
OTHER SOURCE(S): MARPAT 147:46112

AB The present invention relates to a novel compound (e.g., 24-ethyl-cholestane-3β,5α,6α-triol), its production, its use, and to methods of treating neoplasms and other tumors as well as other diseases including hypercholesterolemia, autoimmune diseases, viral diseases (e.g., hepatitis B, hepatitis C, or HIV), and diabetes.

IT 123318-82-1, Clofarabine
RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(treatment of cancer and other diseases using ethylcholestane triol and combination with other agents)

RN 123318-82-1 CAPLUS
CN 9H-Purin-6-amine, 2-chloro-9-(2-deoxy-2-fluoro-β-D-arabinofuranosyl)- (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2004:290469 CAPLUS
DOCUMENT NUMBER: 140:309397
TITLE: Methods and compositions for the treatment of lupus using clofarabine
INVENTOR(S): Wood, Christopher B.; Smith, Stuart William Gordon
PATENT ASSIGNEE(S): Bioenvision, Inc., USA
SOURCE: PCT Int. Appl., 33 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004028463	A2	20040408	WO 2003-US30407	20030925
WO 2004028463	A3	20040715		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2500091	A1	20040408	CA 2003-2500091	20030925
AU 2003276987	A1	20040419	AU 2003-276987	20030925
EP 1551386	A2	20050713	EP 2003-798756	20030925
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
NO 2005002039	A	20050623	NO 2005-2039	20050426
US 20060135463	A1	20060622	US 2005-529520	20051116

PRIORITY APPLN. INFO.:			
		US 2002-414685P	P 20020927
		WO 2003-US30407	W 20030925

AB The invention relates to methods of treating or preventing lupus comprising the administration of clofarabine or a pharmaceutically acceptable salt, hydrate, solvate or clathrate thereof to a patient in need of such treatment. The invention further relates to methods of treating or preventing lupus comprising the administration of clofarabine or a pharmaceutically acceptable salt, hydrate, solvate or clathrate thereof and an addnl. therapeutic agent to a patient in need of

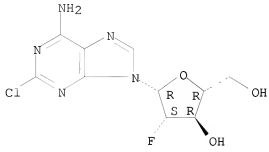
such treatment.

IT 123318-82-1, Clofarabine
RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(methods and compns. for treatment of lupus using clofarabine)

RN 123318-82-1 CAPLUS

CN 9H-Purin-6-amine, 2-chloro-9-(2-deoxy-2-fluoro-β-D-arabinofuranosyl)- (CA INDEX NAME)

Absolute stereochemistry.



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